Claims

- 1. A copolymer comprising the following monomers: acrylic acid or an ester thereof in the range 40 to 80 % by weight; methacrylic acid or an ester thereof in the range 20 to 60 % by weight; and a polymerizable surfactant in the range 0.01 to 9 % by weight.
- 2. A copolymer according to claim 1 comprising the following monomers: ethyl acrylate in the range 40 to 80 % by weight; methyl methacrylate in the range 20 to 60 % by weight; and a monomer characterized by formula I:

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wherein m is an integer from 1-55,
R1 is hydrogen or methyl, and
R2 is hydrogen or a carbon chain having 1 to 20 carbon atoms in the range 0.01 to 9 % by weight.

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- 3. An aqueous polymer dispersion obtainable by polymerization of the following monomers in water in the presence of an emulsifying agent: acrylic acid or an ester thereof in the range 40 to 80 % by weight; methacrylic acid or an ester thereof in the range 20 to 60 % by weight; and a polymerizable surfactant in the range 0.01 to 9 % by weight.
- 4. An aqueous polymer dispersion obtainable by polymerization of the following monomers in water in the presence of an emulsifying agent:

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ethyl acrylate in the range 40 to 80 % by weight;
methyl methacrylate in the range 20 to 60 % by weight;
and a monomer of formula I as described in claim1 in the range 0.01 to 9 % by weight.

- 5. An aqueous polymer dispersion obtainable by polymerization of the following monomers in water in the presence of an emulsifying agent:
 acrylic acid or an ester thereof in the range 40 to 80 % by weight;
 methacrylic acid or an ester thereof in the range 20 to 60 % by weight; and
 a polymerizable surfactant in the range 0.01 to 9 % by weight;
 wherein if the emulsifying agent is an emulsifier with a molecular weight lower than 15 kD then it is partially or fully removed after the polymerization reaction.
 - 6. An aqueous polymer dispersion obtainable by polymerization of the following monomers in water in the presence of an emulsifying agent: ethyl acrylate in the range 40 to 80 % by weight; methyl methacrylate in the range 20 to 60 % by weight; and a monomer of formula I as described in claim 1 in the range 0.01 to 9 % by weight; wherein if the emulsifying agent is an emulsifier with a molecular weight lower than 15 kD then it is partially or fully removed after the polymerization reaction.
 - 7. An aqueous polymer dispersion obtainable by the polymerization of the following monomers in water:
 acrylic acid or an ester thereof in the range 40 to 80 % by weight;
 methacrylic acid or an ester thereof in the range 20 to 60 % by weight; and
 a polymerizable surfactant in the range 0.01 to 9 % by weight.
 - 8. An aqueous polymer dispersion obtainable by the polymerization of the following monomers in water:

ethyl acrylate in the range 40 to 80 % by weight; methyl methacrylate in the range 20 to 60 % by weight; and a monomer of formula I in the range 0.01 to 9 % by weight.

- 9. A film for use in coating pharmaceutical formulations obtainable by removal of water from an aqueous dispersion according to any one of claims 3 to 8.
 - 10. A pharmaceutical formulation comprising:
 - a) a pharmaceutical core comprising a pharmacologically active ingredient; and
- b) a film coating comprising a film according to claim 9.
 - 11. A pharmaceutical formulation comprising a pharmacologically active ingredient which is provided in a plurality of beads wherein each of the beads is coated with a film according to claim 9.
 - 12. A formulation according to either claim 10 or claim 11 wherein the formulation is a controlled release formulation.
- 13. A formulation according to any one of claims 10-12 wherein the pharmacologically active ingredient has activity in the treatment of cardiovascular or gastrointestinal diseases.
 - 14. A formulation according to any one claim 10-12 in which the pharmacologically active ingredient is a beta-blocking adrenergic agent.
- 15. A formulation according to claim 14 in which the pharmacologically active ingredient is metoprolol or a pharmaceutically acceptable salt thereof.
 - 16. A formulation according to claim 15 in which the metoprolol salt is the tartrate, succinate, fumarate or benzoate salt.

17. A polymer according to either claim 1 or claim 2, or a dispersion according to any one of claims 3 to 8 or a film according to claim 9, or a pharmaceutical formulation according to any one of claims 10 to 16 wherein m is 2-55 in the monomer of formula I.

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18. A polymer according to either claim 1 or claim 2, or a dispersion according to any one of claims 3 to 8 or a film according to claim 9, or a pharmaceutical formulation according to any one of claims 10 to 16 wherein the monomer of formula I is defined as m is 4, R1 is hydrogen and R2 has 13 carbon atoms.

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19. A polymer according to either claim 1 or claim 2, or a dispersion according to any one of claims 3 to 8 or a film according to claim 9, or a pharmaceutical formulation according to any one of claims 10 to 16 wherein the monomer of formula I is defined as m is 10, R1 is hydrogen and R2 has 11 carbon atoms.

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20. A polymer according to either claim 1 or claim 2, or a dispersion according to any one of claims 3 to 8 or a film according to claim 9, or a pharmaceutical formulation according to any one of claims 10 to 16 wherein the monomer of formula I is defined as m is 25, R1 is hydrogen and R2 has 18 carbon atoms.

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21. A polymer according to either claim 1 or claim 2, or a dispersion according to any one of claims 3 to 8 or a film according to claim 9, or a pharmaceutical formulation according to any one of claims 10 to 16 wherein the monomer of formula I is defined as m is 1, R1 is methyl and R2 is hydrogen.

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22. A polymer according to either claim 1 or claim 2, or a dispersion according to any one of claims 3 to 8 or a film according to claim 9, or a pharmaceutical formulation according to any one of claims 10 to 16 wherein the monomer of formula I is defined as m is 9, R1 is methyl and R2 is hydrogen.

23. A process for the preparation of a polymer comprising polymerizing the following monomers in water in the presence of an emulsifier:

ethyl acrylate in the range 40 to 80 % by weight;

methyl methacrylate in the range 20 to 60 % by weight.; and

a monomer characterized by formula I:

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wherein m is an integer from 1-55,

R1 is hydrogen or methyl, and

R2 is hydrogen or a carbon chain having 1 to 20 carbon atoms in the range 0.01 to 9 % by weight

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- 24. A process according to claim 23 wherein the process is carried out at a temperature in the range of 1 to 100°C.
- 25. A process to prepare a formulation as claimed in any one of claims 10 to 16
 comprising coating the pharmaceutical core or beads with a film coating dispersion as defined in any one of claims 3 to 8.